(19) World Intellectual Property Organization International Bureau



T (THE THE SHEET IN THE STATE AND A SECURE THE STATE AND A STATE

(43) International Publication Date 7 June 2001 (07.06.2001)

PCT

(10) International Publication Number WO 01/40403 A1

(51) International Patent Classification⁷: C09K 11/79

(21) International Application Number: PCT/EP00/12046

(22) International Filing Date:

30 November 2000 (30.11.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

99123747.0

30 November 1999 (30.11.1999) EF

(71) Applicant (for all designated States except US):
PATENT-TREUHAND-GESELLSCHAFT FÜR
ELEKTRISCHE GLÜHLAMPEN MBH [DE/DE];
Hellabrunner Str. 1, 81543 München (DE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): BRAUNE, Bert [DE/DE]; Gerhardstr. 146, 66333 Voelklingen (DE).
 WAITL, Guenther [DE/DE]; Praschweg 3, 93042
 Regensburg (DE). BOGNER, Georg [DE/DE]; Am Sandbuegel 12, 93138 Hainsacker (DE). HINTZEN,

H.T. [NL/NL]; Laboratory of Solid State and Materials Chemistry, Eindhoven University of Technology, P.O. Box 513 STO 2.26, NL-5600 MB Eindhoven (NL). VAN KREVEL, J.W.H. [NL/NL]; Laboratory of Solid State and Materials Chemistry, Eindhoven University of Technology, P.O. Box 513 STO 2.41, NL-5600 MB Eindhoven (NL). BOTTY, Ir.G. [NL/NL]; Oce Technologies B.V., Dept. DPS-EP1, P.O. Box 101, NL-5900 MA Venlo (NL).

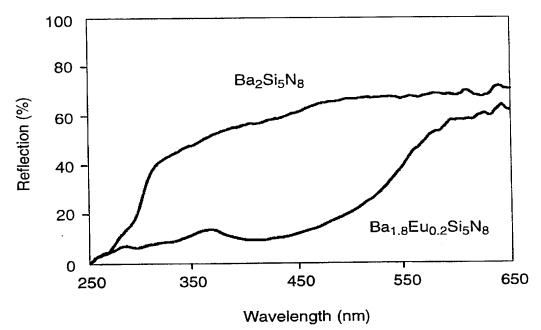
- (74) Common Representative: PATENT-TREU-HAND-GESELLSCHAFT FÜR ELEKTRISCHE GLÜHLAMPEN MBH; Postfach 22 16 34, 80506 München (DE).
- (81) Designated States (national): CA, CN, HU, JP, KR, US.
- (84) Designated States (regional): European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR).

Published:

With international search report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: LIGHT SOURCE USING A YELLOW-TO-RED-EMITTING PHOSPHOR



(57) Abstract: A light source uses a yellow to red emitting phosphor with a host lattice of the nitridosilicate type $M_xSi_yN_z$: Eu, wherein M is at least one of an alkaline earth metal chosen from the group Ca, Sr, Ba and wherein z = 2/3x + 4/3y.

Light source using a yellow-to-red-emitting phosphor

5

10

15

20

Technical Field

This invention relates to an electrical Light source using a yellow-to-redemitting phosphor and more particularly, but not exclusively to Light Emitting Diodes (LED). The phosphor belongs to the class of rare-earth activated silicon nitrides. Other application fields are electrical lamps, especially high intensity discharge lamps or fluorescent lamps.

Background Art

For Eu²⁺ -doped material normally UV-blue emission is observed (Blasse and Grabmeier: Luminescent Materials, Springer Verlag, Heidelberg, 1994). Several studies show that also emission in the green and yellow part of the visible spectrum is possible (Blasse: Special Cases of divalent lanthanide emission, Eur. J. Solid State Inorg. Chem. 33 (1996), p. 175; Poort, Blokpoel and Blasse: Luminescence of Eu²⁺ in Barium and Strontium Aluminate and Gallate, Chem. Mater. 7 (1995), p. 1547; Poort, Reijnhoudt, van der Kuip, and Blasse: Luminescence of Eu²⁺ in Silicate host lattices with Alkaline earth ions in a row, J. Alloys and Comp. 241 (1996), p. 75). Hitherto, red Eu²⁺ luminescence is observed only in some exceptional cases, such as in alkaline earth sulphides and related lattices of the rock-salt type (Nakao, Luminescence centers of MgS, CaS and CaSe Phosphors Activated with Eu²⁺ Ion, J. Phys. Soc. Jpn. 48(1980), p. 534), in alkaline earth thiogallates (Davolos, Garcia, Fouassier, and Hagenmuller, Luminescence of Eu²⁺ in Strontium and Barium Thiogallates, J. Solid. State Chem. 83 (1989), p. 316) and in some borates (Diaz and Keszler; Red, Green, and Blue Eu²⁺ luminescence in solid state Borates: a structure-property relationship, Mater. Res. Bull. 31 (1996),

10

15

20

25

p. 147). Eu²⁺ luminescence in alkaline-earth silicon nitrides has hitherto only been reported for MgSiN₂:Eu (Gaido, Dubrovskii, and Zykov: Photoluminescence of MgSiN₂ Activated by Europium, Izv. Akad. Nauk SSSR, Neorg. Mater. 10 (1974), p. 564; Dubrovskii, Zykov and Chernovets: Luminescence of rare earth Activated MgSiN₂, Izv. Akad. Nauk SSSR, Neorg. Mater. 17 (1981), p. 1421) and Mg_{1-x}Zn_xSiN₂:Eu (Lim, Lee, Chang: Photoluminescence Characterization of Mg_{1-x}Zn_xSiN₂:Tb for Thin Film Electroluminescent Devices Application, Inorganic and Organic Electroluminescence, Berlin, Wissenschaft und Technik Verlag, (1996), p. 363). For both Eu²⁺ luminescence in the green and green/blue part of the spectrum was found.

New host lattices of the nitridosilicate type are based on a three dimensional network of cross-linked SiN₄ tetrahedra in which alkaline earth ions (M= Ca, Sr and Ba) are incorporated. Such lattices are for example $Ca_2Si_5N_8$ (Schlieper and Schlick: Nitridosilicate I, Hochtemperatursynthese und Kristallstruktur von $Ca_2Si_5N_8$, Z. anorg. allg. Chem. 621, (1995), p. 1037), $Sr_2Si_5N_8$ and $Ba_2Si_5N_8$ (Schlieper, Millus and Schlick: Nitridosilicate II, Hochtemperatursynthesen und Kristallstrukturen von $Sr_2Si_5N_8$ and $Ba_2Si_5N_8$, Z. anorg. allg. Chem. 621, (1995), p. 1380), and $BaSi_7N_{10}$ (Huppertz and Schnick: Edge-Sharing SiN₄ tetrahedra in the highly condensed Nitridosilicate $BaSi_7N_{10}$, Chem. Eur. J. 3 (1997), p. 249). The lattice types are mentioned in Table 1.

Sulfide based phosphors (e.g. earth alkaline sulfides) are less desirable for lighting applications, especially for LED applications, because they interact with the encapsulating resin system, and partially suffer from hydrolytic attack. Red emitting Eu²⁺ activated borates show already temperature quenching to a certain degree at the operating temperature of LEDs.

Disclosure of the Invention

It is, therefore, an object of this invention to obviate the disadvantages of the prior art. It is another object of the invention to provide a light source with im-

- 3 -

proved red color rendition R9. It is a further object to provide a light source with an improved overall color rendition Ra. It is a further object to provide a white LED with high color rendition.

Especially high stability up to at least 100 °C is desirable for LED applications. Their typical operation temperature is around 80 °C.

5

10

15

20

25

These objects are accomplished by the characterising features of claim 1. Advantageous embodiments can be found in the dependant claims.

The light source uses a new yellowish-red emitting phosphor. Its absorption is at least within the blue to green spectral region. Furthermore they show fluorescent emission under absorption. Those Eu²⁺-doped luminescent materials show emission within the yellow to red spectral region, especially long wavelength red, orange or yellow emission. These phosphors are based on alkaline-earth silicon nitride material as host-lattices. They are very promising, especially for LED applications, when used as phosphors. Hitherto white LEDs were realised by combining a blue emitting diode with a yellow emitting phosphor. Such a combination has only a poor colour rendition. A far better performance can be achieved by using a multicolor (for example red-greenblue) system. Typically the new material can be used together with a greenemitting (or yellow-emitting) phosphor, for example strontiumaluminate SrAl₂O₄:Eu²⁺, whose emission maximum is around 520 nm.

In detail, the new Light source using a yellow-to-red-emitting phosphor, uses a host lattice of the nitridosilicate type $M_xSi_yN_z$:Eu, wherein M is at least one of an alkaline earth metal chosen from the group Ca, Sr, Ba and wherein z=2/3x+4/3y. The incorporation of nitrogen increases the proportion of covalent bond and ligand-field splitting. As a consequence this leads to a pronounced shift of excitation and emission bands to longer wavelengths in comparison to oxide lattices.

10

Preferably, the phosphor is of the type, wherein x = 2, and y = 5. In another preferred embodiment, the phosphor is of the type, wherein x = 1, and y = 7.

-4-

Preferably, the metal M in the phosphor is strontium because the resulting phosphor is emitting at relatively short yellow to red wavelengths. Thus the efficiency is rather high in comparison to most of the other elected metals M.

In a further embodiment the phosphor uses a mixture of different metals, for example Ca (10 atom.-%) together with Ba (balance), as component M.

These materials show high absorption and good excitation in the UV and blue visible spectrum (up to more than 450 nm), high quantum efficiency and low temperature quenching up to 100 °C.

It can be used for luminescence conversion LEDs with a blue light emitting primary source together with one or more phosphors (red and green). Another field of application are compact fluorescent lamps and replacement of yttrium vanadate in high intensity discharge lamps.

Brief Description of the Drawings

- Fig. 1: Diffuse reflection spectra of undoped Ba₂Si₅N₈ and Ba₂Si₅N₈:Eu;
 - Fig. 2: Diffuse reflection spectra of undoped $BaSi_7N_{10}$ and $BaSi_7N_{10}$:Eu;
 - Fig. 3: Emission spectrum of Ba₂Si₅N₈:Eu;
 - Fig. 4: Emission spectrum of BaSi₇N₁₀:Eu;
 - Fig. 5-7: Emission spectrum of several embodiments of $Sr_2Si_5N_8$:Eu;
- 20 Fig. 8: Emission spectrum of Ca₂Si₅N₈:Eu;
 - Fig. 9: Emission spectra of a white LEDs;
 - Fig. 10: shows a semiconductor element which serves as light source for white light.

- 5 -

Detailed Embodiments

Eu₂O₃ (with purity 99,99 %), or Eu metal (99,99 %), Ba metal (> 99 %); Sr metal (99 %), Ca₃N₂ (98 %), or Ca powder (99,5%) and Si₃N₄ (99,9 %) were used as commercially available starting materials. Ba and Sr were nitrided by firing at 550 and 800 °C under a nitrogen atmosphere. Subsequently, Ca₃N₂ or nitrided Ba, Ca or Sr were ground in a mortar and stoichiometrically mixed with Si₃N₄ under nitrogen atmosphere. The Eu-concentration was 10 atom.% compared to the alkaline earth ion. The powdered mixture was fired in molybdenum crucibles at about 1300-1400 °C in a horizontal tube furnace under nitrogen/hydrogen atmosphere. After firing, the powders were characterised by powder X-ray diffraction (Cu, Kα-line), which showed that all compounds had formed.

5

10

15

20

25

The undoped $Ba_2Si_5N_8$, $Ca_2Si_5N_8$ and $BaSi_7N_{10}$ are greyish-white powders. These undoped rare-earth activated silicon nitrides show high reflection in the visible range (400-650 nm) and a strong drop in the reflection between 250-300 nm (Fig. 1 and 2). The drop in reflectance is ascribed to host-lattice absorption. The Eu-doped samples are orange-red, except for $BaSi_7N_{10}$:Eu which is orange-yellow (Table 1). The strong coloration is unique for Eu^{2+} -doped rare-earth activated silicon nitrides and make these material interesting orange-red phosphors. A typical example of a reflection spectrum of $Ba_2Si_5N_8$:Eu shows that the absorption due to Eu is superposed on the host-lattice absorption and extends up to 500-550 nm (Fig. 1). This explains the red-orange colour of these compounds. Similar reflection spectra were observed for $Sr_2Si_5N_8$:Eu and $Ca_2Si_5N_8$:Eu.

For $BaSi_7N_{10}$:Eu the absorption of Eu is less far in the visible part (Fig. 2), which explains the orange-yellow colour of this compound.

All samples show efficient luminescence under UV excitation with emission maxima in the orange-red part of the visible spectrum (see Table 1). Two typical examples of emission spectra can be seen in Figs. 3 and 4. They

10

show that the emission is at extremely long wavelengths (for Eu²⁺ emission) with maxima up to 660 nm for BaSi₇N₁₀:Eu (Fig. 4.). Excitation bands are observed at low energy which is the result of a centre of gravity of the Eu²⁺ 5d band at low energy and a strong ligand-field splitting of the Eu²⁺ 5d band, as can be expected for N³⁻ containing lattices (van Krevel, Hintzen, Metselaar, and Meijerink: Long Wavelength Ce³⁺-luminescence in Y-Si-O-N Materials, J. Alloys and Comp. 168 (1998) 272).

Since these materials can convert blue into red light due to low-energy excitation bands, they can be applied in white light sources, for example based on primarily blue-emitting LED's (typically GaN or InGaN) combined with red, vellow and/or green emitting phosphors.

Table 1:

Compound	Crystal structure	Colour	Emission Maximum (nm)*
Ca ₂ Si ₅ N ₈ :Eu	Monoclinic	Orange-Red	600 to 630
Sr ₂ Si ₅ N ₈ :Eu	Orthorhombic	Orange-Red	610 to 650
Ba ₂ Si ₅ N ₈ :Eu	Orthorhombic	Orange-Red	620 to 660
BaSi ₇ N ₁₀ :Eu	Monoclinic	Orange-Yellow	640 to 680

*depending on the conditions for preparation and concentration of the activator; typical values for Eu-concentration may vary between 1 and 10% compared to the alkaline-earth ion M

These emission maxima are unusually far in the long wavelength side. A specific example is a phosphor of the type $Sr_{1.8}Eu_{0.2}Si_5N_8$. Its emission spectrum is shown in fig. 5.

Another embodiment for realising M is the use of Zn. It can replace Ba, Sr or Ca fully or partially.

20 A further embodiment for replacing Si fully or partially is Ge. An concrete embodiment is $Sr_{1.8}Eu_{0.2}Ge_5N_8$.

Some further specific examples were investigated.

The preparation conditions and optical properties of the red emitting phosphor $Sr_2Si_5N_8$: Eu^{2+} were investigated. Optimisation showed a quantum efficiency of about 70 %. The emission is tuneable between 610 and 650 nm, depending on the Eu^{2+} concentration in the sample and the heating conditions. The absorption at 400 nm and 460 nm is high (reflection of only 15-40 %) and the temperature quenching of the luminescence at 80° C is low (only 4 %). The particle size of the phosphor is without milling below 5 μ m. These properties make this phosphor very interesting especially for application in both the UV and blue LED.

For the nitride synthesis, the starting materials are Si₃N₄ (99,9% (mainly α-phase),
Alfa Aesar), Sr metal (dendritic pieces 99,9 %, Alfa Aesar) and Eu₂O₃ (4N). The Sr
metal has to be nitrided and in case one uses instead of Eu₂O₃ Eu metal, this has
also to be nitrided.

The Sr metal is milled by hand in an agath mortar in an argon glovebox and nitrided at 800° C under N_2 . This results in a nitration over 80° .

After remilling, the nitrided metal, together with Si₃N₄ and Eu₂O₃, is milled and mixed by hand again in the glovebox. The heating of this mixture has typically the following parameters:

18°C/min to 800°C
5h at 800°C
20 18°C/min to T_{end} (1300-1575°C)
5h at T_{end} (1300-1575°C)
H₂(3.75%)/N₂ 400l/h

 $\text{Ca}_2\text{Si}_5\text{N}_8\text{:Eu}^{2+}$ samples were made with Ca_3N_2 as starting material.

An overview of all the samples is given in table 1. Typically, the samples were first heated at 800°C, and then they were heated a second time in the same cycle at elevated (1300-1600°C) temperatures. The samples were then milled (mill under air), sieved and measured.

10

Table 1: parameters of heating cycles of (Ca,Sr)₂Si₅N₈:Eu²⁺ samples

Code	Ca/Sr	Eu ²⁺ (%)	Time 1 (h)	Temp. 1 (°C)	Time 2 (h)	Temp. 2 (°C)
EC/HU 31/00	Са	10	5	800	5	1400
EC/HU 42/00	Ca	1	5	800	5	1565
EC/HU 41/00	Ca0.4Sr1.4	10	5	800	5	1565
EC/HU 62/00	Sr	1	5	800	5	1400
EC/HU 63/00	Sr	2	5	800	5	1400
EC/HU 64/00	Sr	3	5	800	5	1400
EC/HU 65/00	Sr	5	5	800	5	1400
EC/HU 66/00	Sr	8	5	800	5	1400
EC/HU 67/00	Sr	10	5	800	5	1400

The samples that are obtained after this heating show a color of deep orange for $10\%~Eu^{2+}$ containing $Sr_2Si_5N_8$ samples. With less Eu^{2+} the colour is fainter. The Ca samples have a yellow-orange colour.

There is also another interesting feature: the powder particles are very small with an average particle size d_{50} between about 0,5 and 5 μ m, a typically value is d_{50} =1.3 μ m. The small particle sizes are advantageous for the processing of LEDs with luminescent material. For example they allow a homogeneous distribution in the resin.

Table 2: Optical data of (Ca,Sr)₂Si₅N₈:Eu²⁺ samples

Code	Ca/Sr	Eu ²⁺ (%)	Em. Max (nm)	Refl. 400 (%)	Refl. 460 (%)	QE (%)	×	У
EC/HU 31/00	Ca	10	619	12	19	26	0.600	0.396
EC/HU 42/00	Ca	1	603	47	58	37	0.555	0.435
EC/HU 41/00	Ca0.4 Sr1.4	10	660	17	22	59	0.636	0.,354
EC/HU 62/00	Sr	1	609	53	58	70	0.602	0.393
EC/Hu 63/00	Sr	2	618	43	48	73	0.615	0.381
EC/Hu 64/00	Sr	3	621	36	41	72	0.622	0.374
EC/Hu 65/00	Sr	5	624	26	32	67	0.632	0.365
EC/HU 66/00	Sr	8	636	21	26	67	0.641	0.356
EC/HU 67/00	Sr	10	644	17	22	64	0.642	0.354

10

15

20

25

30

Concerning table 2 all samples were typically first heated in a first cycle (for example 800°C for 5h), as already outlined above.

Included in table 2 are the position of the emission maximum, the mean wavelength, the reflection at 400 and 460 nm, the quantum efficiency and finally the x and y colour coordinates.

From table 2 it can be derived that the pure Ca samples are not as favourable as the Sr samples. It is surprising that the Sr-Ca compound has an emission wavelength that is larger than that of the pure Sr compound.

Specific examples are shown in Figures 6 to 8. Figure 6 shows the energy distribution (in arbitrary units) and reflection (in percent) of sample HU 64/00 ($Sr_2Si_5N_8:Eu^{2+}$) having a proportion of 3% Eu and a quantum efficiency of 72%. Figure 7 shows the energy distribution (in arbitrary units) and reflection (in percent) of sample HU 65/00 ($Sr_2Si_5N_8:Eu^{2+}$) having a proportion of 5% Eu and a quantum efficiency of 67%. Figure 8 shows the energy distribution (in arbitrary units) and reflection (in percent) of sample HU 42/00 ($Ca_2Si_5N_8:Eu^{2+}$) having a proportion of 1% Eu and a quantum efficiency of 37%.

A specific example of a light source is a white LED whose principle is well-known, see for example US-A 5 998 925 or US-A 6 066 861. It uses a blue emitting primary light source (peak emission around 380 to 470 nm), preferably a (In)GaN-chip whose radiation is partially absorbed by a conversion material based on a resin and at least one phosphor. Improved color rendition can be obtained for a white LED using two or three conversion phosphors for a blue light of the primary light source. Embodiments may contain first phosphors with main emission in the range 480 to 600 nm, for example yttrium based garnets, thiogallates and/or chlorosilicates together with second phosphors of the inventive new type rare-earth activated silicon nitrides having a main emission above 600 nm and preferably above 650 nm. Some overlap between the emission of the different phosphors may exist between 600 and 650 nm. Especially by using a white LED based on primary emission between 420 and 470 nm (peak wavelength) together with a first phosphor YAG:Ce (yellow) and a second phosphor M₂Si₅N₈:Eu²⁺ (red) a color

10

15

20

25

30

rendition Ra up to 85 and even around 90 was obtained, depending of details of the mixture and choice of M.

- 10 -

Even further improving is obtained by first phosphors with more greenish emission (maximum around 490 nm).

Another concept is a UV-radiation emitting primary light source for exciting three phosphors (RGB-concept), wherein the Red component is a phosphor of the inventive new type rare-earth activated silicon nitrides and the Green and Blue component are well known phosphors as outlined above.

Besides, the inventive new phosphor can be used to create a highly stable red or orange or yellow emitting LED which may be based on a primary light source (preferably InGaN-Chip) of peak emission around 380 to 480 nm whose light is fully converted by a nitride phosphor of the inventive type rare-earth activated silicon nitrides doped with Eu. These LEDs show higher efficiency and improved stability compared to well-known commercial LEDs with direct excitation of yellow to red colors.

Figure 9 shows the spectral emission of several white LEDs. They are based on an InGaN-Chip emitting primary radiation with a peak emission wavelength of 460 nm which is partially converted within an epoxy resin covering the chip.

The schematic construction of a light source for white light is explicitly shown in Fig. 10. The light source is a semiconductor element (chip 1) of the type InGaN having a peak emission wavelength of 460 nm and a first and a second electrical terminal 2, 3 including a bonding wire 14, the semiconductor element being embedded in the region of a recess 9 within a basic housing 8, which is not transmissive for light. The frontier between housing 8 and recess 9 is a wall which serves as a reflector for the blue primary radiation emitted from chip 1. The recess 9 is filled with a potting mass 5 which comprises as main constituents an epoxy casting resin and phosphor pigments 6 (less than 15% by weight of the mass 5). Further small amounts are provided, among others, by methyl ether and aerosil.

The resin of several embodiments comprises different phosphor compositions. In detail, a reference is given by a standard white LED based on conversion of

- 11 -

the primary emitted blue light only by YAG:Ce phosphor. Its proportion is 3,6 weight-% of the resin (curve 1). Three embodiments comprising $Sr_2Si_5N_8:Eu^{2+}$ in addition to YAG:Ce were investigated. Keeping the whole proportion of conversion phosphors constant (3,6 %), an amount of 0,25 % YAG:Ce was replaced by $Sr_2Si_5N_8:Eu^{2+}$, see curve 2. This amount of $Sr_2Si_5N_8:Eu^{2+}$ was further increased to 0,5 % (curve 3) and to 0,75% (curve 4). Their color rendition Ra was markedly improved by 6% (curve 2), 10% (curve 3), and 12% (curve 4) relative to the reference (curve 1).

Claims

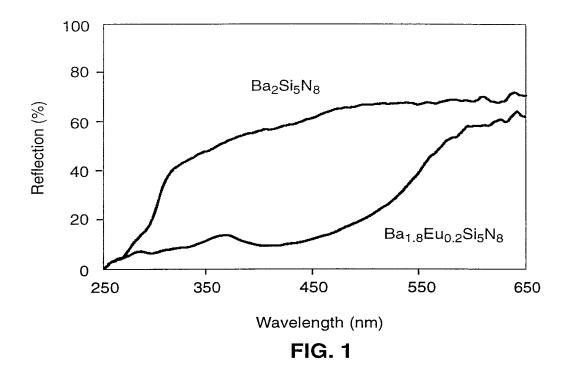
- 12 -

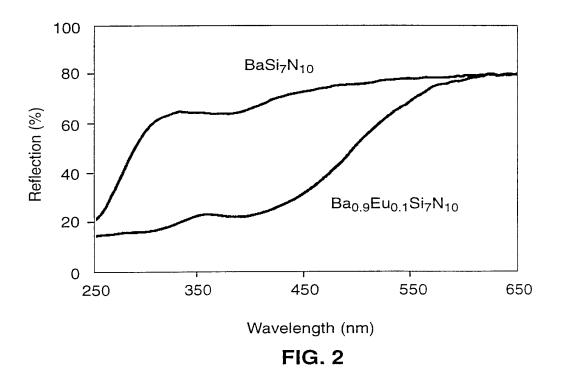
- 1. Light source using a yellow-to-red-emitting phosphor, wherein the phosphor converts at least partially radiation of a primary light source, characterised in that said phosphor has a host lattice of the nitridosilicate type $M_xSi_yN_z$:Eu, wherein M is at least one of an alkaline earth metal chosen from the group Ca, Sr, Ba, Zn and wherein z = 2/3x + 4/3y.
- 2. Light source according to claim 1, wherein x = 2, and y = 5.

5

20

- 3. Light source according to claim 1, wherein x = 1, and y = 7.
- 4. Light source according to claim 1, wherein M is strontium.
- 5. Light source according to claim 1, wherein M is a mixture of at least two metals of said group.
 - 6. Light source according to claim 1, wherein Si is replaced fully or partially by Ge.
 - 7. Light source according to claim 1 wherein the average particle size of said phosphor is between 0,5 and 5 μm .
- 15 8. Light source of claim 1 wherein the light source emits primary radiation and secondary radiation of said phosphor is emitted under excitation by the primary radiation.
 - 9. Light source of claim 8 wherein the primary radiation is blue, preferably between 420 and 470 nm (peak wavelength), and combines with the secondary radiation and optionally with further secondary radiation of at least one further phosphor in order to obtain white light.
 - 10. Light source according to claim 9 with a color rendition index Ra of at least 85, preferably 90.





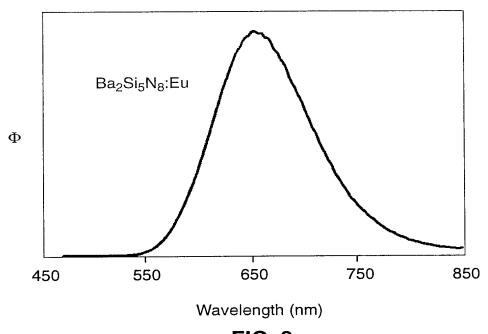


FIG. 3

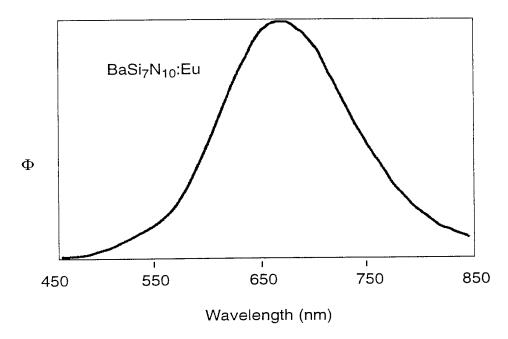
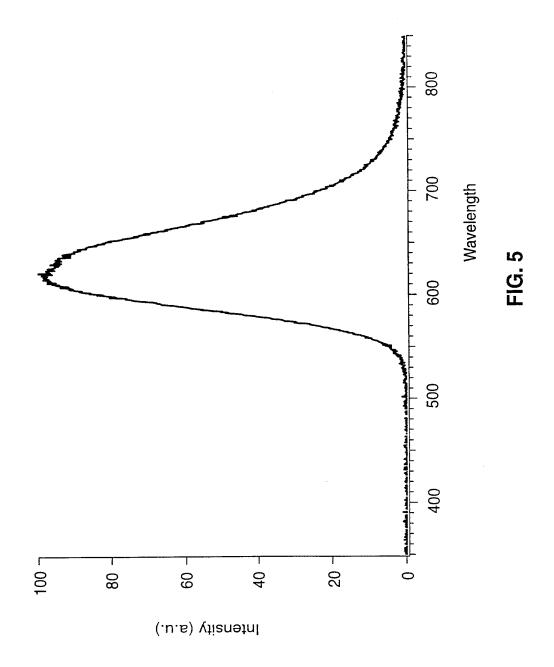
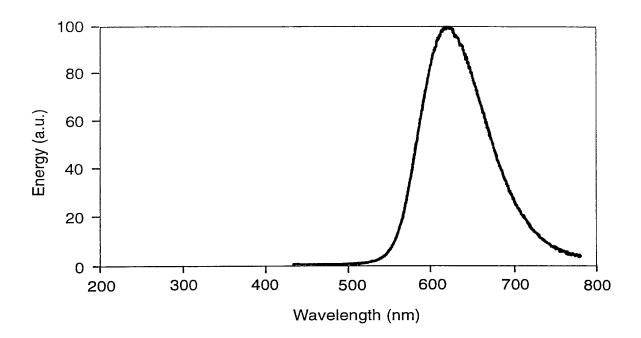


FIG. 4





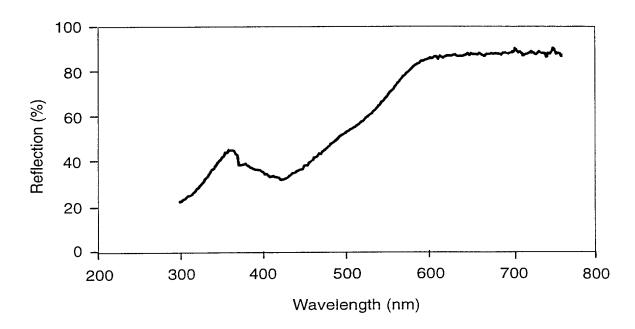
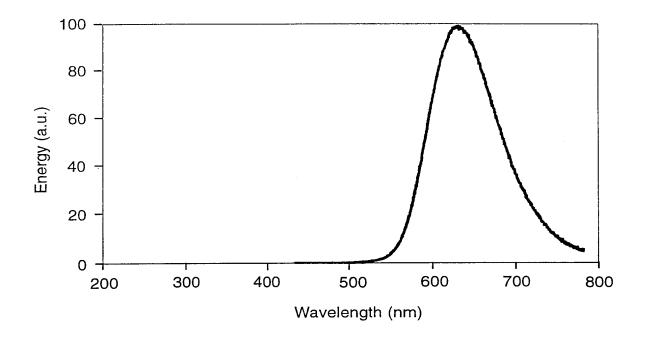


FIG. 6



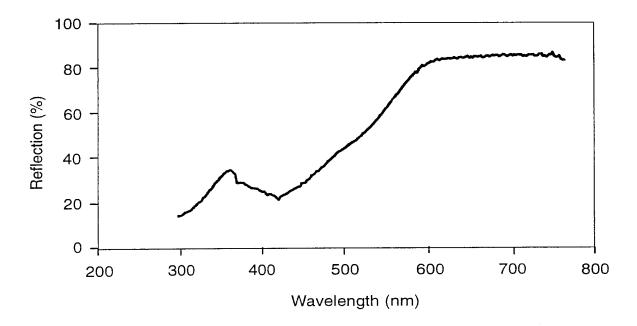
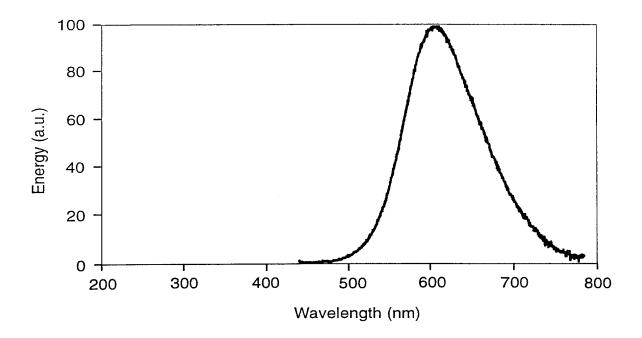


FIG. 7



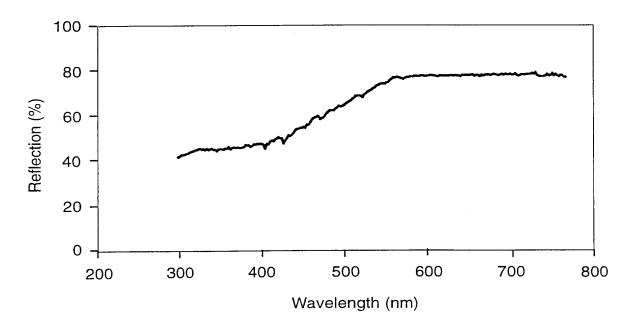
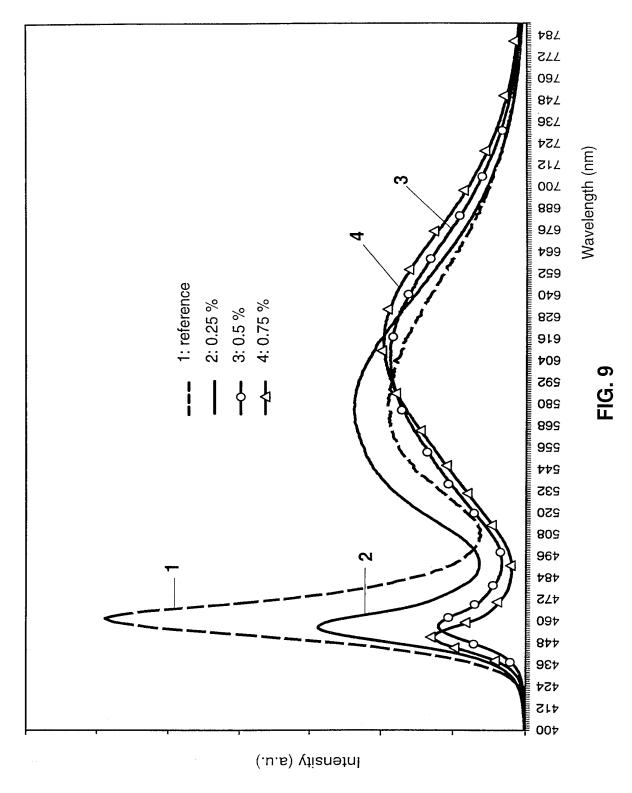


FIG. 8



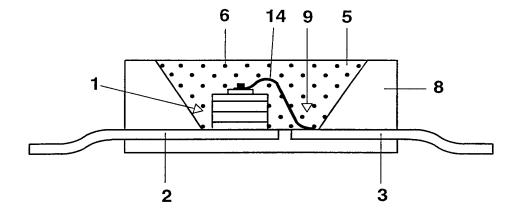


FIG. 10

INTERNATIONAL SEARCH REPORT

Int tional Application No PCT/EP 00/12046

	101/21 00/12010			
A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C09K11/79				
According to International Patent Classification (IPC) or to both national classif	ication and IPC			
B. FIELDS SEARCHED				
Minimum documentation searched (classification system followed by classification $7 - 609$ K	ation symbols)			
Documentation searched other than minimum documentation to the extent that				
Electronic data base consulted during the international search (name of data to INSPEC, WPI Data, PAJ, EPO-Internal, IBM-				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category ° Citation of document, with indication, where appropriate, of the r	relevant passages Relevant to claim No.			
A SOON-SEOK LEE ET AL: "Developme luminescent characteristics of 0 2/ based phosphors" JOURNAL OF THE INSTITUTE OF ELECTION ENGINEERS OF KOREA D, OCT. 1999 ELECTRON. ENG. KOREA, SOUTH KORE vol. 36-D, no. 10, pages 31-36 XP002136109 ISSN: 1226-5845 page 31	CaSiN/sub CTRONICS , INST. EA,			
Further documents are listed in the continuation of box C.	Patent family members are listed in annex.			
Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	 "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family 			
Date of the actual completion of the international search	Date of mailing of the international search report			
12 March 2001	19/03/2001			
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Drouot-Onillon, M-C			